& Clinton St. between 6th & 9th Sts., Arkadelphia, 11000464

COLORADO

Arapahoe County

Englewood Post Office, 3332 S. Broadway, Englewood, 11000465

DISTRICT OF COLUMBIA

District of Columbia

Linnaean Hill (Boundary Increase), 3545 Williamsburg Ln., NW., Washington, 11000466

MINNESOTA

Fillmore County

Bridge No. 5722, N. Section St. over Spring Valley Cr., Spring Valley, 11000467

Houston County

Bridge No. 6679, MN 76 over S. Fork of Root R. (Sheldon Township), Houston, 11000468

Swift County

Gethsemane Episcopal Church, 40 N. Hering St., Appleton, 11000469

Traverse County

District No. 44 School, U.S. 75 (Taylor Township), Campbell, 11000470

MISSISSIPPI

Lee County

Baldwyn Historic District, Roughly along E. & W. Main Sts. & N. & S. 2nd Ave., Baldwyn, 11000471

Webster County

Eupora Historic District, Roughly along N. Dunn St. & W. Roane Ave., Eupora, 11000472

Winston County

Downtown Louisville Historic District, Bounded by Church St., W. Park St., Columbus Ave. & Mill St., Louisville, 11000473

Yazoo County

Rosedale Plantation, 5302 Bend Rd., Vaughan, 11000474

NEW MEXICO

McKinley County

Borrego Pass Trading Post Historic District, Bldg. 1601, Co. Rd. 19, Borrego Pass, 11000475

WASHINGTON

Whatcom County

Lynden Department Store, 444 Front St., Lynden, 11000476

WISCONSIN

Rock County

Eager, Almeron, Funerary Monument and Plot, 8012 N. Cemetery Rd., Evansville, 11000477

Sauk County

Rest Haven Motel, E5116 U.S. 14, Spring Green, 11000478

Vernon County

Vernon County Normal School, 410 S. Center Ave., Viroqua, 11000479

[FR Doc. 2011–16248 Filed 6–28–11; 8:45 am]

BILLING CODE 4312–51–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to § 1301.33(a), Title 21 of the Code of Federal Regulations (CFR), this is notice that on March 14, 2011, Pharmagra Labs, Inc., 158 McLean Road, Brevard, North Carolina 28712, made application to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Pentobarbital (2270), a basic class of controlled substance listed in schedule II.

The company plans to manufacture the listed substances for analytical research and clinical trials.

Any other such applicant, and any person who is presently registered with DEA to manufacture such substances, may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

Any such comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 8701 Morrissette Drive, Springfield, Virginia 22152; and must be filed no later than August 29, 2011.

Dated: June 22, 2011.

Joseph T. Rannazzisi,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 2011–16294 Filed 6–28–11; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By Notice dated April 11, 2011, and published in the **Federal Register** on April 19, 2011, 76 FR 21915, Meda Pharmaceuticals, Inc., 705 Eldorado Street, Decatur, Illinois 62523, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of Nabilone (7379), a basic class of controlled substance listed in schedule II.

The company plans to import the listed controlled substance as a finished drug product in dosage form only for distribution to its customers. The company does not import the listed controlled substance in bulk active pharmaceutical ingredient (API) form.

There are no domestic sources of Nabilone in finished drug product form available in the United States. The U.S. Food and Drug Administration has approved this product for medical use in the United States.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and § 952(a), and determined that the registration of Meda Pharmaceuticals Inc. to import the basic class of controlled substance is consistent with the public interest, and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. DEA has investigated Meda Pharmaceuticals Inc. to ensure that the company's registration is consistent with the public interest. The investigation has included inspection and testing of the company's physical security systems, verification of the company's compliance with state and local laws, and a review of the company's background and history. Therefore, pursuant to 21 U.S.C. 952(a) and 958(a), and in accordance with 21 CFR 1301.34, the above named company is granted registration as an importer of the basic class of controlled substance listed.

Dated: June 22, 2011.

Joseph T. Rannazzisi,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 2011–16286 Filed 6–28–11; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Office of Justice Programs

[OJP; BJA Docket No. 1561]

Meeting of the Department of Justice's (DOJ's) National Motor Vehicle Title Information System (NMVTIS) Federal Advisory Committee

AGENCY: Bureau of Justice Assistance, Justice.

ACTION: Notice of meeting.

SUMMARY: This is an announcement of a meeting of DOJ's National Motor Vehicle Title Information System (NMVTIS) Federal Advisory Committee to discuss the role of the NMVTIS Federal Advisory Committee Members and various issues relating to the operation and implementation of NMVTIS.